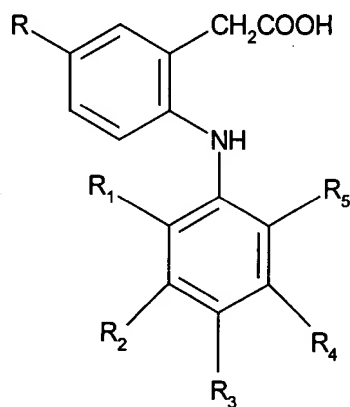


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wherein R is methyl or ethyl;

R<sub>1</sub> is chloro or fluoro;

R<sub>2</sub> is hydrogen or fluoro;

R<sub>3</sub> is hydrogen, fluoro, chloro, methyl, ethyl, methoxy or ethoxy;

R<sub>4</sub> is hydrogen or fluoro;

R<sub>5</sub> is chloro, fluoro or trifluoromethyl;

or a pharmaceutically acceptable salt thereof;

or a pharmaceutically acceptable prodrug ester thereof.

18 17  
38. A method according to claim 37 wherein the compound is a compound of formula I wherein R is methyl or ethyl; R<sub>1</sub> is chloro or fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen, fluoro, chloro or methyl; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro or fluoro; or a pharmaceutically acceptable salt thereof; or a pharmaceutically acceptable prodrug ester thereof.

19 17  
39. A method according to claim 37 wherein the compound is a compound of formula I wherein R is methyl or ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen or fluoro; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; or a pharmaceutically acceptable salt thereof; or a pharmaceutically acceptable prodrug ester thereof.

30  
40. A method according to claim 37 wherein the compound is a compound of formula I wherein R is methyl or ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is fluoro; R<sub>3</sub> is hydrogen or ethoxy; R<sub>4</sub> is fluoro; and R<sub>5</sub> is fluoro; or a pharmaceutically acceptable salt thereof; or a pharmaceutically acceptable prodrug ester thereof.

21  
41. A method according to claim 37 wherein the compound is a compound of formula I wherein R is methyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen or fluoro; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; or a pharmaceutically acceptable salt thereof; or a pharmaceutically acceptable prodrug ester thereof.

22  
42. A method according to claim 37 wherein the compound is a compound of formula I wherein R is methyl or ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub>-R<sub>4</sub> are hydrogen or fluoro; and R<sub>5</sub> is chloro or fluoro; or a pharmaceutically acceptable salt thereof; or a pharmaceutically acceptable prodrug ester thereof.

23  
43. A method according to claim 37 wherein the compound is 5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid of formula I wherein R is methyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; or a pharmaceutically acceptable salt thereof.

24  
44. A method according to claim 37 wherein the compound is 5-methyl-2-(2',4'-difluoro-6'-chloroanilino)phenylacetic acid of formula I wherein R is methyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is fluoro; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; or a pharmaceutically acceptable salt thereof.

25  
45. A method according to claim 37 wherein the compound is 5-ethyl-2-(2',3',5',6'-tetrafluoroanilino)phenylacetic acid of formula I wherein R is ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is fluoro; R<sub>3</sub> is hydrogen; R<sub>4</sub> is fluoro; and R<sub>5</sub> is fluoro; or a pharmaceutically acceptable salt thereof.

#### REMARKS

Reconsideration of the application as amended is respectfully requested.

The claims under consideration in the Office Action were claims 1-8, 10-13 and 15-35. New claim 36 was apparently inadvertently omitted from the Office Action summary. Claims 20-35 stand allowed. Claims 1-8, 10-13, 15-19 stand rejected over DE 3,445,011 (Ciba) and US 3,958,690 (Sallmann, *et al.*). EP 865,788 (Yamazaki, *et al.*) was withdrawn as a reference against the claims as amended.